old In-Tube Test QuantiFERON®-TB G

Mycobacterium tuberculosis infection. This test was approved by the U.S. Food and Drug Administration (FDA) in 2007. The QuantiFERON®-TB Gold In-Tube test (QFT-IT) is a serological whole-blood test for use as an aid in diagnosing This test cannot, in and of itself, rule in or out active tuberculosis disease.

WHAT ARE THE ADVANTAGES?

- Requires a single patient visit to draw blood sample.
- Results can be available within 24 hours.
- Does not boost responses measured by subsequent tests.
- Is not subject to reader bias.
- Is not affected by prior BCG (bacille Calmette-Guérin) vaccination.
- immunosuppressive therapy). Those unable to mount an immune response will most likely produce an indeterminate tes Can alert providers to patients with impaired T-cell immunity (e.g., persons with HIV, cancer, renal failure, or undergoing

WHAT ARE THE DISADVANTAGES AND LIMITATIONS?

- Blood samples must arrive at the Utah Public Health Laboratory within 15 hours of collection, or be incubated and spun within strict guidelines (contact UPHL for more information on time constraints)
- acquired immunodeficiency syndrome [AIDS], current treatment with immunosuppressive drugs, selected hematological There is limited data on the use of QFT-IT in children younger than 17 years of age, among persons recently exposed disorders, specific malignancies, diabetes, silicosis, chronic renal failure, and in pregnant women). Refer to the most to M. tuberculosis, and in immunocompromised persons (e.g., impaired immune function caused by HIV infection or natter. recent publications available for current information on this rr
 - Errors in collecting or transporting blood specimens or in running and interpreting the assay can decrease the accuracy of QFT-IT.
- for developing TB disease. Limited data on the use of QFT-IT to determine who is at risk

QuantiFERON®-TB Gold In-Tube testing is available at the Utah Public Health Laboratory. For more information please

Although not available at the Utah Public Health Laboratory, there is another interferon gamma release assay (IGRA) tesi available called the T-Spot $^{ ext{ iny B}}$ which was FDA approved on July 25, 200 call (801) 584-8452 or visit: http://www.health.utah.gov/lab

HEALTH DEPARTMENTS **DIRECTORY OF UTAH LOCAL**

BEAR RIVER

Phone: (435) 792-6500 655 East 1300 North Fax: (435) 792-6600 Logan, Utah 84341

Room 209

Salt Lake City, Utah 84111 Phone: (801) 534-4600 610 South 200 East, Fax: (801) 534-4557

SOUTHEASTERN UTAH 28 South 100 East

Phone: (435) 637-3671 Fax: (435) 637-1933 Price, Utah 84501 PO Box 800

Phone: (435) 896-5451 Richfield, Utah 84701

70 Westview Drive

CENTRAL UTAH

Fax: (435) 896-4353

620 South 400 East, SOUTHWEST UTAH

50 East State St. - Basement

PO Box 618

Courthouse Annex

DAVIS COUNTY

St. George, Utah 84770 Phone: (435) 986-2551 Fax: (435) 652-4069 Suite 400 Farmington, Utah 84025 Phone: (801) 451-3393

Fax: (801) 451-3464

Department of Health

SALT LAKE VALLEY

Phone: (801) 851-7029 151 South University Fax: (801) 343-8729 Avenue, Suite 1800 Provo, Utah 84601

Kamas, Utah 84036 Phone: (435) 783-4351 110 North Main Street

Fax: (435) 783-6021

WASATCH COUNTY

Heber City, Utah 84032 Phone: (435) 654-2700 Fax: (435) 654-2705 55 South 500 East

Tooele, Utah 84074 Phone: (435) 277-2310 Fax: (435) 277-2314

151 North Main Street

OELE COUNTY

NEBER-MORGAN

133 South 500 East

TRI COUNTY

Vernal, Utah 84078

Phone: (801) 399-7252 Fax: (801) 399-7260 Ogden, Utah 84401 477 23rd Street

Phone: (435) 247-1177 Fax: (435) 781-0536

area (see listing on back cover). and/or to the local health department in your Utah Department of Health Report all suspected and confirmed cases to the

288 North 1460 West Box 142105 Salt Lake City, Utah 84114-2105 Phone: (801) 538-6993 Fax: (801) 538-9913

TB Control/Refugee Health Program Utah Department of Health



Post-Test Referral

Guidelines for

Testing for TB Infection

REKCIPOZIZ



TUBERCULOSIS IS A REPORTABLE DISEASE

Fold Line

Required by Utah State Statute

Provider Guide to Testing for TB Infection and Post-Test Referral

Step 1

Classify the Results for TB Infection Test²

The following measurements of induration are classified as positive in a Tuberculin Skin Test (TST):^{\phi}

QuantiFERON® (QFT®):Ω

QuantiFERON® Positive

HIV-positive persons

Persons with evidence of old, healed and untreated tuberculosis (TB) on a chest x-ray

≥ 5 mm

- Recent contacts of persons with active TB disease
- Patients with organ transplants and other immunocompromised patients
- Persons with medical risk factors for TB (Table 2)

≥ 10 mm

- Substance abusers
- Recent arrivals from high incidence areas (Table 1)
- Persons at higher risk for exposure to or infection with TB (Table 1)
- Mycobacteriology lab personnel
- Children under age 5
- Children/adolescents exposed to adults in high-risk categories (Tables 1 & 2)

Persons at low risk for TB disease for whom testing is not generally indicated

≥ 15 mm

Mycobacterium
tuberculosis (MTB)
infection likely in most
circumstances.
Refer questions to the
Utah Public Health
Laboratory at
(801) 584-8452

Step 2

If indicated, obtain a chest x-ray and a medical evaluation

Any person with a newly positive TST or QuantiFERON® test result, including high-priority contacts of a patient with active TB disease (as defined in Step 1), should have a chest x-ray to evaluate for active TB disease. If the initial chest x-ray is normal, no follow-up chest x-rays are indicated.

Step 3

Are TB symptoms present, or is the chest x-ray abnormal?

Yes

- Fever
- Chills
- Fatigue
- Loss of appetite
- Weight loss
- Night sweats
- Prolonged productive cough
- Chest pain
- Coughing up blood



Evaluate for active TB disease

Refer for treatment according to the guidelines in Table 3

Table 1. Persons at higher risk for exposure to or infection with TB

- Close contacts of persons known or suspected to have active TB disease.
- Foreign-born persons from areas where TB is common. See: www.health.utah.gov/cdc/tb_home.htm to view up-to-date *M. tuberculosis* incidence information by country.
- Residents and employees of high-risk congregate settings (e.g., correctional institutions, nursing homes, mental institutions, other long term residential settings, homeless shelters).
- Health care workers who serve high-risk patients.
- Medically underserved, low income populations.
 High-risk racial or ethnic minority
- populations.

 Children exposed to adults in high-risk
- categories.Persons who inject illicit drugs.

Table 2. Medical risk factors for the development of active TB disease in TB-infected patients

 HIV infection (or risk for HIV in patients who decline HIV testing).

New TB infection within the previous

- two years.

 Evidence of old, healed TB on a chest x-rav.
- x-ray.Diabetes.
- End-stage renal disease.
- Prolonged corticosteroid therapy.
- Other immunosuppressive therapy.Cancer of the head and neck.
- Hematologic and reticuloendothelial diseases (e.g., leukemia and Hodgkin's disease).
- Silicosis.
- Chronic malabsorption syndromes.
- Intestinal bypass or gastrectomy.Being 10% or more below ideal body
- weight.
 Substance abuse.

Table 3. Guidelines for treatment of latent tuberculosis infection by patient risk factors, TST result, QuantiFERON® result, and age* ¥

CANDIDATES FOR TREATMENT OF LATENT TUBERCULOSIS INFECTION (LTBI)

CATEGORY OF PERSON TESTED	TST < 5 mm	TST ≥ 5 mm	TST ≥ 10 mm	TST ≥ 15 mm	or	QFT®-Pos ^π	QFT®-Neg
Case Contact: Children < age 5*	Treat**	Treat	Treat	Treat		Treat	Treat**
Case Contact: HIV-infected [©]	Treat**	Treat	Treat	Treat		Treat	Treat**
Case Contact: Immunocompromised*	Treat**	Treat	Treat	Treat		Treat	Treat**
Case Contact: ≥ age 5 and immunocompetent $^{\Sigma}$	Repeat***	Treat	Treat	Treat		Treat	Repeat***
Immunocompromised persons	Do Not Treat	Treat	Treat	Treat		Treat	Do Not Treat
HIV-infected	Do Not Treat	Treat	Treat	Treat		Treat	Do Not Treat
Fibrotic changes on chest x-ray	Do Not Treat	Treat	Treat	Treat		Treat	Do Not Treat
Recent arrival from endemic country $^{\Phi}$	Do Not Treat	Do Not Treat	Treat	Treat		Treat	Do Not Treat
Injection drug user	Do Not Treat	Do Not Treat	Treat	Treat		Treat	Do Not Treat
Resident/employee in an institutional setting§	Do Not Treat	Do Not Treat	Treat	Treat		Treat	Do Not Treat
Mycobacteriology lab personnel	Do Not Treat	Do Not Treat	Treat	Treat		Treat	Do Not Treat
High-risk clinical conditions [‡]	Do Not Treat	Do Not Treat	Treat	Treat		Treat	Do Not Treat
Child < age 5	Do Not Treat	Do Not Treat	Treat	Treat		Treat	Do Not Treat
Persons < age 17 exposed to high-risk adults	Do Not Treat	Do Not Treat	Treat	Treat		Treat	Do Not Treat
No risk factors (TST discouraged)	Do Not Treat	Do Not Treat	Do Not Treat	Treat		Treat	Do Not Treat

For more information on treatment or to refer a client for treatment, please see the Directory of Local Health Departments (back cover).

- £ The window period is the time span between the date of an initial tuberculin skin test (TST) or QuantiFERON® (QFT®) with a negative reaction and the date of the follow-up TST or QFT® that should take place 8-10 weeks after exposure. After the window period has ended, a repeat test should be administered to each contact who had an initial negative reaction.
- * QFT® is not FDA approved to test those under 5 years of age. Case contacts who are under 5 years of age or immunocompromised, initially testing negative should be started on therapy. Testing should be repeated 8-10 weeks after last exposure to TB. Treatment can be discontinued after second negative TST or QFT® in children. Immunocompromised patients with a second negative TST or QFT® need to be evaluated by a physician.
- ** Treat and repeat TST or QFT® test after 8-10 weeks.
- *** Repeat TST or QFT® test after 8-10 weeks.
- HIV-infected contacts should receive a full course of treatment even if they have a second reaction of < 5 mm or a negative QFT®.
- ¥ TST and QFT® are not contraindicated for persons who have been vaccinated with BCG (bacille Calmette-Guérin). Test results for *M. tuberculosis* infection for individuals with a history of BCG should be interpreted by using the same diagnostic cut points used for individuals without a history of BCG vaccination.
- π Rarely QFT[®] may cross-react with *M. kansasii*, *M. szulgai*, or *M. marinum* resulting in a false-positive result.
- π Harely QF1° may cross-react with *M. Kansasii, M. Szuigai, or M. marinum* resulting in a false-positive result.

 Φ See *www.health.utah.gov/cdc/tb home.htm* to view up-to-date *M.tuberculosis* incidence information by country.
- § TST Conversion: An increase in reaction size of ≥ 10 mm within 2 years should be considered a TST conversion indicative of recent infection with *M. tuberculosis*.
- ‡ Silicosis, diabetes mellitus, chronic renal failure, some hematologic disorders (e.g. leukemias and lymphomas), other specific malignancies (e.g. carcinoma of the head and neck or lung), being ≥ 10% below ideal body weight, gastrectomy, intestinal bypass.
- In all situations of high suspicion of tuberculosis, a person with a negative TST or QFT® should be further evaluated with radiological, bacteriological, HIV, or other

Tuberculin Skin Test - Results are classified according to a patient's risk factor for tuberculosis. In general, a history of vaccination with BCG (bacille Calmette-Guerin) does not influence the need for tuberculin skin testing, the classification of TST results, or clinical decisions regarding the management of TST-positive individuals. See Tables 1, 2, and 3.

QuantiFERON® - Negative indicates infection unlikely, but cannot be excluded if symptoms are consistent with MTB disease, if test is less than 8-10 weeks after exposure, or individual has increased likelihood of disease progression. See disadvantages and limitations on back cover.

^Ψ**High-Priority Contacts** - Certain high-priority contacts need medical follow-up even if their reaction is less than 5 mm because they are at high risk of both developing active TB disease and having a false-negative TST result. These include: (1) immunocompromised contacts and (2) children younger than age 5 who were tested less than 8-10 weeks after the last exposure to TB. No further evaluation is necessary when high-priority contacts have a negative reaction to a TST given *more* than 8 weeks after the last exposure to TB. **See Step 2 and Table 3 footnote (*).**